Preliminary Amendment Filed September 8, 2003

Amendments to the Claims

Cancel claims 1-26 and insert new claim 27 as shown below.

Claims 1-26 (Cancelled)

Claim 27 (New) A method for treating diabetic retinopathy in a mammal in need of such treatment comprising administering an effective $\alpha_v \beta_3$ inhibiting amount comprising from about 0.01 mg to about 1000 mg per kilogram of body weight of a compound of the formula

$$A \xrightarrow{\left(\begin{array}{c} Y^3 \\ Z^3 \end{array}\right)_t} B \xrightarrow{\left(\begin{array}{c} Q \\ S \end{array}\right)_t} COR$$

or a pharmaceutically acceptable salt thereof, wherein

B is selected from the group consisting of -CONR⁵⁰- and -SO₂NR⁵⁰-;

wherein Y¹ is selected from the group consisting of N-R², O, and S;

R² is selected from the group consisting of H; alkyl; aryl; hydroxy; alkoxy; cyano; nitro; amino; alkenyl; alkynyl; alkyl optionally substituted with one or more substitutent selected from lower alkyl, halogen, hydroxyl, haloalkyl, cyano, nitro, carboxyl, amino, alkoxy, aryl or aryl optionally substituted with one or more halogen, haloalkyl, lower alkyl, alkoxy, cyano, alkylsulfonyl, alkylthio, nitro, carboxyl, amino, hydroxyl, sulfonic acid, sulfonamide, aryl, fused aryl,

monocyclic heterocycles, or fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, hydroxy, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, cyano, nitro, alkylsulfonyl, sulfonic acid, sulfonamide, carboxyl derivatives, amino, aryl, fused aryl, monocyclic heterocycles and fused monocyclic heterocycle; monocyclic heterocycles; and moncyclic heterocycles optionally substituted with one or more substitutent selected from halogen, haloalkyl, lower alkyl, alkoxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, aryl or fused aryl; or

R² taken together with R⁷ forms a 4-12 membered dinitrogen containing heterocycle optionally substituted with one or more substituent selected from the group consisting of lower alkyl, hydroxy and phenyl; or

R² taken together with R⁷ forms a 5 membered heteroaromatic ring; or

R² taken together with R⁷ forms a 5 membered heteroaromatic ring fused with a phenyl group;

R⁷ (when not taken together with R²) and R⁸ are independently selected from the group consisting of H; alkyl; alkenyl; alkynyl; aralkyl; cycloalkyl; bicycloalkyl; aryl; acyl; benzoyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxy, haloalkyl, cyano, nitro, carboxyl derivatives, amino, alkoxy, thio, alkylthio, sulfonyl, aryl, aralkyl, aryl optionally

substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethyl, sulfonyl, alkylsulfonyl, haloalkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, moncyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, aryloxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, aryl, fused aryl; monocyclic and bicyclic heterocyclicalkyls; -SO₂R¹⁰- wherein R¹⁰ is selected from the group consisting of alkyl, aryl and monocyclic heterocycles, all optionally substituted with one or more substituent selected from the group consisting of halogen, haloalkyl alkyl, alkoxy, cyano, nitro, amino, acylamino, trifluoroalkyl, amido, alkylaminosulfonyl, alkylsufonyl, alkylsulfonylamino, alkylamino, dialkylamino, trifluoromethylthio, trifluoroalkoxy, trifluoromethylsulfonyl, aryl, aryloxy, thio, alkylthio, and monocyclic heterocycles; and

wherein R¹⁰ is defined above; or

NR⁷ and R⁸ taken together form a 4-12 membered mononitrogen containing moncyclic or bicyclic ring optionally substituted with one or more substitutent selected from lower alkyl, carboxyl derivatives, aryl or hydroxy and wherein said

Preliminary Amendment Filed September 8, 2003

ring optionally contains a heteroatom selected from the group consisting of O, N, and S;

R⁵ is selected from the group consisting of H, alkyl, alkenyl, alkynyl, benzyl, and phenethyl; or

wherein Y² is selected from the group consisting of alkyl; cycloalkyl; bicycloalkyl; aryl; monocyclic heterocycles; alkyl optionally substituted with aryl which can also be optionally substituted with one or more substitutent selected from halo, haloalkyl, alkyl, nitro, hydroxy, alkoxy, aryloxy, aryl, or fused aryl; aryl optionally substituted with one or more substituent selected from halo, haloalkyl, hydroxy, alkoxy, aryloxy, aryl, fused aryl, nitro, methylenedioxy, ethylenedioxy, or alkyl; alkylnyl; alkenyl; -SR9 and -OR9- wherein R9 is selected from the group consisting of H; alkyl; aralkyl; aryl; alkenyl; and alkynyl; or R9 taken together with R7 forms a 4-12 membered mononitrogen containing sulfur or oxygen containing heterocyclic ring; and

R⁵ and R⁷ are as defined above; or

Y² (when Y² is carbon) taken together with R⁷ forms a 4-12 membered mononitrogen containing ring optionally substituted with alkyl, aryl, or hydroxy;

Z¹, Z², Z⁴, and Z⁵ are independently selected from the group consisting of H; alkyl; hydroxy; alkoxy; aryloxy; arylalkoxy; halogen; haloalkyl; haloalkoxy; nitro; amino; aminoalkyl; alkylamino; dialkylamino; cyano; alkylthio;

Preliminary Amendment Filed September 8, 2003

alkylsulfonyl; carboxyl derivatives; acetamide; aryl; fused aryl; cycloalkyl; thio; monocyclic heterocycles; fused monocyclic heterocycles; and A, wherein A is defined above;

R⁵⁰ is selected from the group consisting of H and alkyl;

R¹ is selected from the group consisting of H, alkyl, alkenyl, alkynyl, aryl and aryl, optionally substituted with one or more substituent selected form the group consisting of halogen, haloalkyl, hydroxy, alkoxy, aryloxy, aralkoxy, amino, aminoalkyl, carboxyl derivatives, cyano and nitro;

t is an integer 0, 1, or 2;

R is X-R³ wherein X is selected from the group consisting of O, S and NR⁴, wherein R³ and R⁴ are independently selected from the group consisting of hydrogen; alkyl; alkenyl; alkynyl; haloalkyl; aryl; arylalkyl; sugars; steroids and in the case of the free acid, all pharmaceutically acceptable salts thereof; and

 Y^3 and Z^3 are independently selected from the group consisting of H, alkyl, aryl, cycloalkyl and aralkyl.